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	Application No.	Applicant(s)
Notice of Allowability	10/748,089	KONRADI ET AL.
Notice of Allowability	Examiner	Art Unit
	Deepak Rao	1624
The MAILING DATE of this communication appears on the cover sheet with the correspondence address All claims being allowable, PROSECUTION ON THE MERITS IS (OR REMAINS) CLOSED in this application. If not included herewith (or previously mailed), a Notice of Allowance (PTOL-85) or other appropriate communication will be mailed in due course. THIS NOTICE OF ALLOWABILITY IS NOT A GRANT OF PATENT RIGHTS. This application is subject to withdrawal from issue at the initiative of the Office or upon petition by the applicant. See 37 CFR 1.313 and MPEP 1308.		
1. This communication is responsive to the amendment filed on June 22, 2005.		
2. The allowed claim(s) is/are 75-83.		
3. The drawings filed on are accepted by the Examiner.		
4. ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) ☐ All b) ☐ Some* c) ☐ None of the: 1. ☐ Certified copies of the priority documents have been received. 2. ☐ Certified copies of the priority documents have been received in Application No 3. ☐ Copies of the certified copies of the priority documents have been received in this national stage application from the International Bureau (PCT Rule 17.2(a)). * Certified copies not received: Applicant has THREE MONTHS FROM THE "MAILING DATE" of this communication to file a reply complying with the requirements noted below. Failure to timely comply will result in ABANDONMENT of this application. THIS THREE-MONTH PERIOD IS NOT EXTENDABLE. 5. ☐ A SUBSTITUTE OATH OR DECLARATION must be submitted. Note the attached EXAMINER'S AMENDMENT or NOTICE OF INFORMAL PATENT APPLICATION (PTO-152) which gives reason(s) why the oath or declaration is deficient. 6. ☐ CORRECTED DRAWINGS (as "replacement sheets") must be submitted. (a) ☐ including changes required by the Notice of Draftsperson's Patent Drawing Review (PTO-948) attached 1) ☐ hereto or 2) ☐ to Paper No./Mail Date (b) ☐ including changes required by the attached Examiner's Amendment / Comment or in the Office action of Paper No./Mail Date Identifying indicia such as the application number (see 37 CFR 1.84(c)) should be written on the drawings in the front (not the back) of each sheet. Replacement sheet(s) should be labeled as such in the header according to 37 CFR 1.121(d). 7. ☐ DEPOSIT OF and/or INFORMATION about the deposit of BIOLOGICAL MATERIAL must be submitted. Note the attached Examiner's comment regarding REQUIREMENT FOR THE DEPOSIT OF BIOLOGICAL MATERIAL.		
Attachment(s) 1. Notice of References Cited (PTO-892) 2. Notice of Draftperson's Patent Drawing Review (PTO-948) 3. Information Disclosure Statements (PTO-1449 or PTO/SB/0 Paper No./Mail Date 4. Examiner's Comment Regarding Requirement for Deposit of Biological Material	6. ☐ Interview Summary Paper No./Mail Dat 8), 7. ☑ Examiner's Amendn	te

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EXAMINER'S AMENDMENT

An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

Authorization for this examiner's amendment was given in a telephone interview with Mr. Stephen Todd on June 28, 2005.

The application has been amended as follows:

In the claims:

In claim 75, page 3, line 3, delete "and R⁸ are" and in place insert -- is --.

In claim 75, page 3, delete lines 10-13 (paragraph containing definition of R¹⁶).

In claim 81, lines 4-5, delete "pharmaceutical composition of Claim 78" and in place insert -- compound of Claim 75 --.

In claim 82, lines 4-5, delete "pharmaceutical composition of Claim 78" and in place insert -- compound of Claim 76 --.

In claim 83, lines 4-5, delete "pharmaceutical composition of Claim 78" and in place insert -- compound of Claim 77 --.

(Copy of claims 75 and 81-83 as amended in enclosed in the Appendix)

The terminal disclaimer filed on June 22, 2005 disclaiming the terminal portion of any patent granted on this application, which would extend beyond the expiration date of U.S. Patent

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No. 6,492,372 has been reviewed and is accepted. The terminal disclaimer has been recorded.

The obviousness-type double patenting rejection is withdrawn in view of the terminal disclaimer.

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Deepak Rao whose telephone number is (571) 272-0672. The examiner can normally be reached on Tuesday-Friday from 6:30am to 5:00pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, James O. Wilson, Acting-SPE of 1624, can be reached at (571) 272-0661. The fax phone number for the organization where this application or proceeding is assigned is (571) 273-8300.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (571) 272-1600.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Primary Examiner
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June 28, 2005

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APPENDIX

Claim 75 (Currently amended): A compound of formula A or B:

wherein

R^{4"} is selected from the group consisting of hydrogen and alkyl;

R⁵ is selected from the group consisting of alkyl, substituted alkyl, alkenyl, substituted alkenyl, aryl, substituted aryl, cycloalkyl, substituted cycloalkyl, cycloalkenyl, substituted cycloalkenyl, heterocyclic, substituted heterocylic, heteroaryl and substituted heteroaryl;

R⁶ is selected from the group consisting of hydrogen, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkenyl, heterocyclic, substituted heterocyclic, aryl, substituted aryl, heteroaryl, substituted heteroaryl, and -SO₂R¹⁰ where R¹⁰ is selected from the group consisting of alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl,

cycloalkenyl, substituted cycloalkenyl, heterocyclic, substituted heterocyclic, aryl, substituted aryl, heteroaryl, substituted heteroaryl;

R⁷ and R⁸ are is independently selected from the group consisting of hydrogen, alkyl,

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substituted alkyl, cycloalkyl, substituted cycloalkyl, aryl, substituted aryl, heteroaryl, substituted heterocyclic and halogen;

R¹¹ and R¹¹ are independently selected from the group consisting of alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, substituted cycloalkenyl, heterocyclic, substituted heterocyclic, and where R¹¹ and R¹¹ are joined to form a heterocycle or a substituted heterocycle;

R¹⁶-is independently selected from the group consisting of hydrogen, alkyl, substitutedalkyl, alkoxy, substituted alkoxy, amino, substituted amino, cycloalkyl, substituted cycloalkyl, aryl, substituted aryl, heterocyclic and halogen; and

R¹⁸ is selected from the group consisting of alkyl, substituted alkyl, alkoxy, substituted alkoxy, amino, substituted amino, cycloalkyl, substituted cycloalkyl, nitro, aryl, substituted aryl, heteroaryl, substituted heteroaryl, heterocyclic and substituted heterocyclic;

R²⁰ is selected from the group consisting of hydrogen, alkyl, substituted alkyl, alkoxy, substituted alkoxy, cycloalkyl, substituted cycloalkyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, heterocyclic, substituted heterocyclic and halogen;

X is selected from the group consisting of hydroxyl, alkoxy, substituted alkoxy, alkenoxy, substituted alkenoxy, cycloalkoxy, substituted cycloalkoxy, cycloalkenoxy, substituted cycloalkenoxy, substituted heteroaryloxy, heteroaryloxy, substituted heteroaryloxy, substituted heterocyclyloxy and—NR"R" where each R" is independently selected from the group consisting of hydrogen, alkyl, substituted alkyl, alkenyl, substituted alkenyl, cycloalkyl, substituted cycloalkyl, aryl, substituted aryl, heteroaryl, substituted heterocyclic;

x is an integer of from 0 to 4.

or enantiomers, diastereomers and pharmaceutically acceptable salts thereof.

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Claim 81 (Currently amended): A method for treating inflammation in a mammalian patient which inflammation is mediated by VLA-4, which method comprises administering to said patient a pharmaceutical composition comprising a pharmaceutically acceptable excipient and a pharmaceutically effective amount of a compound pharmaceutical emposition of Claim 78 75.

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Claim 82 (Currently amended): A method for treating inflammation in a mammalian patient which inflammation is mediated by VLA-4, which method comprises administering to said patient a pharmaceutical composition comprising a pharmaceutically acceptable excipient and a pharmaceutically effective amount of a compound pharmaceutical emposition of Claim 79 76.

Claim 81 (Currently amended): A method for treating inflammation in a mammalian patient which inflammation is mediated by VLA-4, which method comprises administering to said patient a pharmaceutical composition comprising a pharmaceutically acceptable excipient and a pharmaceutically effective amount of a compound pharmaceutical composition of Claim 80 77.